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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/815,490	03/31/2004	Toshihisa Takeyama	KOT-0094	8491
23413 CANTOR COI	7590 09/26 LBURN, LLP .	007	EXAMINER	
55 GRIFFIN R	OAD SOUTH		ANGEBRANNDT, MARTIN J	
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Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

		Application No.	Applicant(s)		
Office Action Summary		10/815,490			
		Examiner	TAKEYAMA, TOSHIHISA		
		,	Art Unit		
	The MAILING DATE of this communication app	Martin J. Angebranndt  ears on the cover sheet with the co	1756 correspondence address		
Period for Reply					
WHIC - Exte after - If NC - Failu Any	ORTENED STATUTORY PERIOD FOR REPLY CHEVER IS LONGER, FROM THE MAILING DAISIONS of time may be available under the provisions of 37 CFR 1.13 SIX (6) MONTHS from the mailing date of this communication. O period for reply is specified above, the maximum statutory period we re to reply within the set or extended period for reply will, by statute, reply received by the Office later than three months after the mailing ed patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATION  36(a). In no event, however, may a reply be tin  will apply and will expire SIX (6) MONTHS from  cause the application to become ABANDONE	N. nely filed the mailing date of this communication. D (35 U.S.C. § 133).		
Status					
1)⊠	Responsive to communication(s) filed on 7/18/0	07 & 6/7/07.			
		action is non-final.			
3)□	Since this application is in condition for allowance except for formal matters, prosecution as to the merits is				
	closed in accordance with the practice under E	x parte Quayle, 1935 C.D. 11, 45	53 O.G. 213.		
Disposiți	on of Claims	•			
5)□ 6)⊠ 7)□	Claim(s) 1 and 3-20 is/are pending in the application of the above claim(s) is/are withdraw Claim(s) is/are allowed.  Claim(s) 1 and 3-20 is/are rejected.  Claim(s) is/are objected to.  Claim(s) are subject to restriction and/or	n from consideration.			
Applicati	on Papers				
10) 🔲 -	The specification is objected to by the Examiner The drawing(s) filed on is/are: a) acce Applicant may not request that any objection to the d Replacement drawing sheet(s) including the correction of the oath or declaration is objected to by the Example 1.	pted or b) objected to by the E rawing(s) be held in abeyance. See on is required if the drawing(s) is obj	e 37 CFR 1.85(a). ected to. See 37 CFR 1.121(d).		
Priority u	nder 35 U.S.C. § 119				
<ul> <li>12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).</li> <li>a) All b) Some * c) None of:</li> <li>1. Certified copies of the priority documents have been received.</li> <li>2. Certified copies of the priority documents have been received in Application No.</li> <li>3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).</li> <li>* See the attached detailed Office action for a list of the certified copies not received.</li> </ul>					
Attachment	(s)				
2) ☐ Notice 3) ☑ Inform	e of References Cited (PTO-892) e of Draftsperson's Patent Drawing Review (PTO-948) eation Disclosure Statement(s) (PTO/SB/08) No(s)/Mail Date <u>6/7/07</u> .	4) Interview Summary ( Paper No(s)/Mail Da 5) Notice of Informal Pa 6) Other:	te		

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1. The response of the applicant has been read and given careful consideration. Responses to the arguments of the applicant are presented after the first rejection to which they are directed. The rejections of the previous office action are withdrawn based upon the amendment to the claims.

- 2. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:
  - (a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.
- 3. Claims 1,4,5,7-9,14,16,17 and 20 are rejected under 35 U.S.C. 103(a) as being unpatentable over Kawabata et al. '340, in view of Kouge et al. '596.

Kawabata et al. '340 in examples 16-18, a mixture of cationically curable materials, free radical curable materials (bis(4-acryloxydiethoxyphenyl)methane), a radically polymerization initiator, a sensitizing dye (dye 1) and a triphenylsulfonium hexafluorophosophate as the cationic polymerization initiators (table 4, col 11) which was applied to a glass plate and overcoated with a polyethylene film, exposed to the interference light and then postcured with a flood exposure from a mercury lamp (7/55-8/40). The use of various onium salts is disclosed. (6/8-18).

Kouge et al. '596 teaches the use of various sulfonium salts for cationically polymerizing materials using light and/or heat (3/12-19). These are evidenced in table 2, 3 and 5 to thermally cure more rapidly than similar triphenylsulfonium salts.

It would have been obvious to one skilled in the art to modify to cited example of Kawabata et al. '340 by using the inventive sulfonium compounds in place of the triarylsulphonium compound used as the photoinitiator with a reasonable expectation of forming

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a useful photosensitive composition based upon the disclosure of equivalence by Kouge et al. '596 and the resulting composition would be thermally cationically curable at a temperature in the 40-150° C.

The binder compounds disclosed at [0148-0157] of the prepub are small molecules, which are capable of forming a binder. They are not polymeric. The claims do not require that these be polymers, so while it might be considered a little misleading, the claims as interpreted through the specification clearly does not require these to be polymers. The examiner is well ware that there are two initiation systems, but the applicant's analysis fails to appreciate that the sulfonium salts can be activated either through heat or light (UV). The applicant also fails to appreciate that sulfonium salts, including those set forth in the claims, inherently can initiate either free radical or cationic polymerization as evidenced by Kawabata et al. '340 and if the applicant wishes to exclude the case where a single compound can be used for either type of initiation (thermal vs. photo or free radical vs cationic), then the photoinitatior should be specified. If the applicant intends to further describe these, then the specification should be amended to identify those compounds disclosed in the references cited at [0095] which of the references they are found in and provide copies of those references. The claims rejected under this heading are directed to the composition, and so the argument regarding use are merely intended use, the exception being claim 20, which does not describe the curing of the binder compound, but does describe irradiating the composition with light or subjecting it to heat. If the intent is to limit the claims to the thermal cure, then the claims should have this as a required step. The applicant argues as if two different polymerization systems are not taught by Kawabata et al. '340, this is flawed on its face. The modification of Kawabata et al. '340

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through the teachings of Kouge et al. '596 is merely exchanging one sulfonium for another, noting that there is a disclosure of equivalence between only seven sulfonium compounds, so there cannot be excessive picking and choosing required.

The rejection has been modified to address the added limitation, with Kouge et al. '596 establishing the ability of the sulfonium salts disclosed in that reference to initiate polymerization at lower temperatures than similar triphenylsulfonium salts. Therefore, the showing in the declaration of Toshihisa TAKEYAMA is not unexpected in view of the prior art and cannot be held to establish unobvious results. Although it is moot, the examiner notes that the comparison was with compounds in column 1, not the identified column 3.

4. Claims 1,3-5,7-9,14,16,17 and 20 are rejected under 35 U.S.C. 103(a) as being unpatentable over Ohkuma et al. '634, in view of Kouge et al. '596.

Ohkuma et al. '634 in examples 6-8, a mixture of cationically curable materials, free radical curable materials, a radically polymerization initiator (4-diethylamino-ethylbenzoate), a sensitizing dye (dye 1, methylene blue) and a triphenylsulfonium trifluoroacetic acid as the cationic polymerization initiators (col 15) which was applied to a glass plate and overcoated with a polyethylene film to a thickness of 7 to 23 microns, exposed to the interference light and then postcured with a flood exposure from a mercury lamp (11/4-65). The use of various onium salts is disclosed. (9/35-10/30).

It would have been obvious to one skilled in the art to modify to cited example of Ohkuma et al. '634 by using the dilakylaryl or cyclic sulfonium compounds in place of the triarylsulphonium compound used as the photoinitiator with a reasonable expectation of forming a useful photosensitive composition based upon the disclosure of equivalence by Kouge et al.

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'596. The trifluoroacetic acid is present as the counter ion and is consider acid multiplying (see prepub of instant specification at [0108]).

5. Claims 1,3-9,14,16,17 and 20 are rejected under 35 U.S.C. 103(a) as being unpatentable over Ohkuma et al. '634, in view of Kouge et al. '596, further in view of Otaki et al. '744

Otaki et al. '744 teach cationically curable materials including epoxies, cyclic ethers and oxetane rings (5/25-42).

In addition to the basis above, it would have been obvious to use other cationically curable moieties, such as oxetane, in place of epoxide rings used in the media resulting from the combination of Ohkuma et al. '634 and Kouge et al. '596 with a reasonable expectation of forming a useful holographic recording medium based upon the disclosure of equivalence by Otaki et al. '744.

In response to applicant's arguments against the references individually, one cannot show nonobviousness by attacking references individually where the rejections are based on combinations of references. See *In re Keller*, 642 F.2d 413, 208 USPQ 871 (CCPA 1981); *In re Merck & Co.*, 800 F.2d 1091, 231 USPQ 375 (Fed. Cir. 1986). The Ohkuma et al. '634 and Otaki et al. '744 references are both within the holographic field and all the references are concerned with photopolymerization, including cationic polymerization and so are analogous. One skilled in the art would expect the cationically curable monomers disclosed by Otaki et al. '744 to be curable in the system resulting form the combination of Ohkuma et al. '634 and Kouge et al. '596 and based upon their use in holography, be suitable to for holographic processing. The rejection stands.

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6. Claims 1,3-5,7-14 and 16-20 are rejected under 35 U.S.C. 103(a) as being unpatentable over Dhar et al. '551, in view of Ohkuma et al. '634, Kouge et al. '596, Roth et al. '814 and Hegel et al. '008.

Dhar et al. '551 teach in example 1, an acrylate monomer and CGI-784 as the photoinatiator mixed with matrix precursors dibutyltin dilaurate, diisocyanate terminated polypropylene glycol and dihydroxypolypropylene glycol, which are heated (13/65-15). Examples 3 and 4 are similar, place the composition between two glass slides with a spacer and after curing of the matrix are used to record holograms. The ability to form thick recording layers of more than 200 microns is disclosed. (3/13-19,4/3-12). Useful photoactive monomers including acrylates are disclosed as useful in this system. (6/51-67). A reduction is shrinkages of the hologram is also realized (7/1-48). The use of various reactions including cationic epoxy or vinyl ether polymerization to form the matrix is disclosed. (6/26-50)

Roth et al. '814 establishes that sulfonium salts are able to thermally initiate cationic polymerization (1/9+).

Hegel et al. '008 teach in example 1, an acrylate monomer and IRG-784 as the photoinatiator mixed with matrix precursors dibutyltin diacetate, diisocyanate terminated polypropylene glycol and dihydroxypolypropylene glycol, which are placed between 1.2 mm substrates with a 500 micron spacer and after curing of the matrix [0031-0040]. The formation of holograms using these is disclosed. [0052-0056]. Useful photoactive monomers including acrylates are disclosed as useful in this system. [0029]. A reduction is shrinkages of the hologram is also realized. The provision of antireflection coatings on one or both of the substrates is disclosed. [0016,0018]. The substrates may be 0.5-1.3 mm thick [0019].

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It would have been obvious to modify the teachings of Dhar et al. '551 by using an epoxy precursor such as those resulting form the combination of Ohkuma et al. '634 and Kouge et al. '596 as discussed above for the matrix based upon the disclosure and the evidence of compatibility in holographic systems including free radically curable compositions from Ohkuma et al. '634 and the evidence from Roth et al. '814 that sulfonium salts are known to act as thermal cationic curing agents and to use the glass substrates, spacers and AR coatings taught by Hegel et al. '008 based upon the similarity with Dhar et al. '551.

The applicant argues that Dhar et al. does not disclose the initiators recited in the claims. This is correct, but the rejection is not based merely upon this reference and the matrix corresponds to the binder of the applicant's claims. Roth et al. teaches the curing at low temperatures as low as 20 degrees C, which is comparable to the room temperature (25 degrees C) described in Dhar et al. '551. Therefore the combination is reasonable to one skilled in the art and self consistent. The scope of the thermal initiators embraced both Kouge et al. '596 and Roth et al. '814 there is no data to support a position of unobvious results over other sulfonium salts.

7. Claims 1,3-5 and 7-20 are rejected under 35 U.S.C. 103(a) as being unpatentable over Dhar et al. '551, in view of Ohkuma et al. '634, Kouge et al. '596, Roth et al. '814 and Hegel et al. '008, further in view of Horimai et al., WO 02/15176

Horimai et al., WO 02/15176 (Hormai et al. '891 is US equivalent) teaches with respect to figure 1, a holographic recording medium which comprises a substrate (2), a holographic recording layer (photopolymers) (3), a second substrate (4) and a reflective layer (5). The reflective layer and the recording layer can be next to each other (12/5-23; 11/40-64). The interference fringes results from the interferences from the light passing through the layer toward

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the reflective layer and that reflected back into the laser from the reflective layer. (col 5. ?; 4/53-

5/7).

In addition to the basis provided above, it would have been obvious to one skilled in the art to modify the embodiments rendered obvious by the combination of Dhar et al. '551 with Ohkuma et al. '634, Kouge et al. '596, Roth et al. '814 and Hegel et al. '008 as set forth above by adding a reflective layer on the further substrate as taught Horimai et al., WO 02/15176 to

allow holographic recording without a second beam.

8. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Martin J. Angebranndt whose telephone number is 571-272-1378.

The examiner can normally be reached Monday- Fridays.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Mark Huff can be reached on 571-272-1385. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 5/71-272-1000.

Martin J Angebranndt

9/20/02